

Food and Drug Administration (FDA)

- Has regulations for manufacturers to follow while researching new chemical entities and developing those chemicals into drug products
- Continues oversight even drugs are approved
- Issues guidelines as to how the product may be packaged, labeled, advertised, and marketed to physicians (and now the public)
- Manufacturers may not make speculative or false claims
- OTC medications also undergo this scrutiny
 - labels of OTC products must conform to a preferred format

Fda Guidelines Reproduction 1966

Roman Wölfel



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